

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (Currently Amended) Method for discovering suitable elution chromatography parameters for the separation of biological molecules, comprising, in an automated manner:
 - a) arranging different chromatography media in a location dependent manner on a multiwell plate defined by columns (X direction) and rows (Y direction) as matrix, on the matrix points of the plate defined by the matrix in the respective cavities therein, where the chromatography media consists on the one hand of materials (B) which bind the biological sample and on the other hand of materials (NB) which do not bind the biological sample,
 - b) bringing the different chromatography media into contact with a biological sample in the respective cavities,
 - c) where the chromatography media are arranged in the individual cavities of the multiwell plate in such a way that on the one hand a chromatography medium from group B and group NB is present in each individual cavity, but on the other hand this chromatography medium from group B and group NB differ at least in a single parameter,
 - d) separating the biological sample located in the respective cavities into biomolecules bound to binding materials and biomolecules not bound to binding materials, and
 - e) analyzing the bound and not-bound molecules of the biological sample for each individual cavity depending on the chromatography medium located in the respective cavity,

whereby suitable elution chromatography parameters for separating the biological sample into biomolecules are ascertained.

2. **(Previously Presented)** Method according to Claim 1, where the biological sample is purified or unpurified proteins, peptides, nucleic acids of all types, carbohydrates, lipids and other biomolecule substance classes or low-molecular-weight metabolism products or mixtures thereof.
3. **(Previously Presented)** Method according to Claim 1, where the chromatography media of the materials binding the biological sample (group B) are selected from solid particles having the property of absorbing biomolecules.
4. **(Previously Presented)** Method according to Claim 1, where the chromatography media of the compounds not binding the biological sample (group N13) are selected from organic and/or inorganic acids, bases, salts, derivatives thereof or solvents of all types, and aqueous solutions thereof.
5. **(Previously Presented)** Method according to Claim 1, where agents for stabilisation of the biological sample contain at least one of: glycerol, sucrose, sodium molybdate, ethylene glycols, urea, guanidinium chloride, betaine, taurine, DTE, DTT, EDTA, EGTA, monothioglycerol, detergents, polyethylene glycol (PEG), chloroform, methanol, H₂O, protease inhibitors or mixtures thereof.
6. **(Previously Presented)** Method according to Claim 1, where the duration of the bringing into contact of the biological sample with the chromatography media can be freely selected.

7. (Cancelled)

8. (Withdrawn) Kit for discovering suitable chromatography conditions in the separation of biological molecules by the method according to claim 1, comprising:
- a) a multiwell plate which is defined by columns (X direction) and rows (Y direction) as matrix, where different chromatography media are arranged in a location-dependent manner on the matrix points of the plate defined by the matrix,
 - b) different chromatography media for stocking the matrix points.
9. (Previously Presented) Kit according to Claim 8, which contains software for the evaluation, identification and interpretation of the method of the invention.
10. (Previously Presented) Method according to claim 3, wherein the chromatography media of the materials binding the biological sample is affinity chromatography media, anion exchanger media, hydrophobic interaction chromatography media, hydroxylapatite chromatography media, cation exchanger media, metal affinity chromatography media, or reversed-phase material media.